

Examination showed that the device consisted of a mono-polar electrode, a single stage amplifier, and a power supply, the output from which was applied to a section of wire mesh attached beneath a sheet of bakelite.

**LIBELED:** 1-2-58, W. Dist. Wis.; amended libel, 4-9-58.

**CHARGE:** 502(a)—the labeling accompanying the article, while held for sale, contained false and misleading representations that the device was capable of measuring nerve interference; of determining accurately the precise line of drive of a subluxation (an incomplete or partial dislocation); of measuring the resistance of a subluxation; of giving the precise line of drive that would most efficiently reduce nerve interference at any given time; of measuring "some types of energy"; of amplifying the impulses which indicate nerve transmission interference; of determining where, when, how much, and how to administer adjustment; and that such diagnostic use of the device could allay, correct, or prevent sickness, disease, death, chronic ailments in children, improper kidney function, and the abnormal function of the stomach and any other organ or part of the body.

**DISPOSITION:** I. N. Toftness, claimant, filed motions for the return of the seized property, the suppression of evidence and dismissal of the action and on 4-9-58, the court denied the motions. Argument was then heard on claimant's objections to answering the written interrogatories which had been served upon him. The court issued an order directing that the claimant answer all of the interrogatories, but that those answers which would "reveal secret processes, development, and information obtained by research all of which was confidential while looking forward to obtaining a patent," could be separated and placed in a sealed envelope subject only to inspection by the court and the Government. The claimant subsequently withdrew his claim, and, on 6-20-58, judgment of condemnation was entered and the devices were ordered delivered to the Food and Drug Administration for display purposes with the provision that they be sold or destroyed after the expiration of one year.

**5775. Filter Queen vacuum cleaner.** (F.D.C. No. 41990. S. No. 20-481 P.)

**QUANTITY:** 28 devices at Kansas City, Mo.

**SHIPPED:** Between 5-1-58 and 7-10-58, from Chicago, Ill., and Cleveland, Ohio, by Health-Mor, Inc.

**LABEL IN PART:** (Top of device) "Filter Queen."

**ACCOMPANYING LABELING:** Folders entitled "Your Doctor Approves Filter Queen Home Sanitation System" and sales manuals entitled "Filter Queen \* \* \* One of the most important developments in home sanitation in 50 years."

**RESULTS OF INVESTIGATION:** The article was a canister-type vacuum cleaner equipped with a cone-shaped cellulose filter intended to remove dirt particles from the inducted air.

**LIBELED:** 8-20-58, W. Dist. Mo.

**CHARGE:** 502(a)—the labeling of the article, when shipped, contained false and misleading representations that the article was capable of preventing streptococcic infection of the sinuses, lungs, brain, spinal cord, blood, joints, middle ear bones, tuberculosis, scarlet fever and diphtheria; preventing such disease conditions as erysipelas, acute abscess and peritonitis, angina, scarlet fever, bronchopneumonia, meningitis, pleurisy, mastoiditis, septic sore throat, arthritis, pulmonary tuberculosis, septic diphtheria, smallpox, measles, an-

thrax, tetanus, whooping cough, enteritis, asthma, and skin and lung cancer; and would protect against such disease-forming bacteria as staphylococci, streptococci, sarcinae, tetrads, hemolytic and non-hemolytic bacteria, and gas formers.

DISPOSITION: 8-21-58. Consent—claimed by Filter Queen of Mid-America, Inc., Kansas City, Mo., and relabeled.

5776. Healthmore chair. (F.D.C. No. 41706. S. No. 28-502 P.)

QUANTITY: 13 devices at New Orleans, La.

SHIPPED: 5-1-58, from North Hollywood, Calif., by Wizard Mfg. Co.

LABEL IN PART: "Healthmore Model 500 \* \* \* Custom Built by Wizard Mfg. Co. North Hollywood Calif."

ACCOMPANYING LABELING: Leaflets entitled "Wizard Healthmore Chair."

RESULTS OF INVESTIGATION: The device was indicated to be an upholstered, foam rubber cushioned, reclining, and oscillating chair equipped with a heating element and capable of providing vibration.

LIBELED: 5-15-58, E. Dist. La.

CHARGE: 502(a)—the labeling of the device, when shipped, contained false and misleading representations that the device was effective for overcoming arthritis, bursitis, rheumatism, poor blood circulation, fatigue, tension, and for providing improved health.

DISPOSITION: 9-12-58. Consent—claimed by Wizard Mfg. Co. and relabeled.

5777. Vibrator device. (F.D.C. No. 42071. S. No. 26-457 P.)

QUANTITY: 14 devices at Des Moines, Iowa.

SHIPPED: At various times, after 1-1-58, from Milwaukee, Wis., by Hallmark System.

LABEL IN PART: (Metal plate on device) "Hallmark System."

ACCOMPANYING LABELING: Brochures entitled "A New Idea By Hallmark."

RESULTS OF INVESTIGATION: The device was an upholstered, carrying case-type device containing an electric motor providing vibration.

LIBELED: 7-31-58, S. Dist. Iowa.

CHARGE: 502(a)—the labeling accompanying the article, when shipped and while held for sale, contained false and misleading representations that the article was an adequate and effective treatment for arthritis, bursitis, rheumatism, relieving simple constipation, stimulating blood circulation, relieving aches and pains, breaking down fatty tissues, and reducing "anywhere you have excess weight."

DISPOSITION: 9-9-58. Default—delivered to Food and Drug Administration.

5778. Uranium Wonderpads and Uranium Wondergloves. (F.D.C. No. 41880. S. No. 9-394 P.)

QUANTITY: 5 *Uranium Wondergloves* and 9 *Uranium Wonderpads* at Buffalo, N.Y.

SHIPPED: 5-26-58, from Jackson, Pa., by Jackson Uranium Corp.

LABEL IN PART: "Uranium Wonderpad [or "Wonderglove"] Jackson Uranium Corp. Susquehanna, Pa."

ACCOMPANYING LABELING: Leaflets entitled "Uranium Wonderglove," and "Uranium Wonderpad"; a form letter entitled "The Jackson Uranium Corpo-